Food and Drug Administration, HHS

§522.2340 Sulfomyxin.

- (a) Specifications. Sulfomyxin for injection is sterile. It is derived from the antibiotic substance produced by the growth of Bacillus polymyxa or is the same substance produced by any other means.
- (b) *Sponsor*. See No. 000069 in §510.600(c) of this chapter.
- (c) Special considerations. The quantities of antibiotic in paragraph (e) of this section refer to the activity of the appropriate standard.
- (d) Related tolerances. See §556.700 of this chapter.
- (e) Conditions of use. (1) It is used or intended for use in chickens and turkeys as an aid in the treatment of disease caused or complicated by E. coli, such as colibacillosis and complicated chronic respiratory disease.
- (2) It is administered by subcutaneous injection as follows:

Age of birds in days	Antibiotic activity		
	Chickens (units)	Turkeys (units)	
1 to 14	12,500 25,000 50,000 50,000	12,500 25,000 50,000 100,000	

- (3) A second injection may be given 3 days later if symptoms persist.
- (4) Not for use in laying hens; do not treat chickens within 5 days of slaughter; do not treat turkeys within 7 days of slaughter.

§ 522.2404 Thialbarbitone sodium for injection.

- (a) Specifications. Thialbarbitone sodium for injection when reconstituted with sterile distilled water provides 94 milligrams of thialbarbitone sodium per milliliter of solution.
- (b) *Sponsor*. See No. 000856 in §510.600(c) of this chapter.
- (c) Conditions of use. (1) The drug is administered as a general anesthetic in surgical procedures on dogs, cats, swine, sheep, cattle, and horses. The drug is used for procedures of relatively short duration. However, the period of anesthesia can be lengthened by slower initial injection and supplemental administration during surgery.
- (2) It is administered intravenously. The drug is injected slowly to dogs, cats, cattle, sheep, and swine. For

horses, it is recommended that a preanesthetic sedation be administered to the horse 30 minutes before the drug is administered. The drug is then injected rapidly and completely. The drug is used at the following dosage levels:

Species	Weight of animal in pounds	Dosage in milligrams per pound
Dog	Over 50	14.1
Do	30–50	18.8
Do	10–30	23.5
Do	Under 10	28.2
Cat		31.3-37.6
Horse		6.3-7.8
Cattle and swine		6.7–9.4
Calves and sheep		9.4–11.8

(3) Federal law restricts this drug to use by or on the order of a licensed veterinarian.

§ 522.2424 Sodium thiamylal for injection.

- (a) Specifications. The drug is a sterile dry powder. It is reconstituted aseptically with sterile distilled water, water for injection, or sodium chloride injection, to a desired concentration of 0.5 to 4 percent sodium thiamylal.
- (b) *Sponsors*. See code Nos. 000010 and 000856 in §510.500(c) of this chapter.
- (c) Conditions of use. (1) It is used as an ultra-short-acting anesthetic in dogs, cats, swine, horses, and cattle.
- (2) When diluted aseptically to the desired concentration and administered intravenously to effect, the average single dose is:
- (i) Dogs and cats: 8 milligrams per pound of body weight (when used with a preanesthetic, generally one-half the normal dose).
- (ii) Swine: 40 milligrams per 5 pounds of body weight.
- (iii) Horses: Light anesthesia, 1 gram per 500 pounds to 1,100 pounds of body weight; deep anethesia, 1 gram per 300 pounds of body weight (40 milligrams per 12 pounds of body weight).
- (iv) Cattle: Short duration, 20 milligrams per 5 pounds of body weight; longer duration, 40 milligrams per 7 pounds of body weight.
- (3) Federal law restricts this drug to use by or on the order of a licensed veterinarian.
- (4) NAS/NRC status: The conditions of use specified in this paragraph are NAS/NRC reviewed and found effective.

§ 522.2444

Applications for these uses need not include effectiveness data as specified in §514.111 of this chapter, but may require bioequivalency and safety information.

[40 FR 25812, June 19, 1975, as amended at 49 FR 8434, Mar. 7, 1984; 53 FR 23390, June 22, 1988; 53 FR 40728, Oct. 18, 1988; 62 FR 35076, June 30, 1997]

§ 522.2444 Sodium thiopental implantation or injectable dosage forms.

§ 522.2444a Sodium thiopental for injection.

- (a) Specifications. The drug contains sodium thiopental sterile powder for dilution with sterile water for injection
- (b) Sponsor. See No. 000856 in §510.600(c) of this chapter.
- (c) Conditions of use. (1) It is used as an anesthetic for intravenous administration to dogs and cats during short to moderately long surgical and other procedures. It is also used to induce anesthesia in dogs and cats which then have surgical anesthesia maintained by use of a volatile anesthetic.
 - (2) It is administered as follows:
- (i) For brief anesthesia (6 to 10 minutes) a dosage of 6 to 9 milligrams per pound of body weight is suggested.
- (ii) To obtain anesthesia of 15 to 25 minutes duration the suggested dosage is 10 to 12 milligrams per pound of body weight.
- (iii) Use of a preanesthetic tranquilizer or morphine will decrease the dosage of sodium thiopental required, provide for smoother induction and smoother recovery, and sometimes prolong the recovery period. If morphine is used as a preanesthetic agent the dose of the barbiturate can be reduced as much as 40 to 50 percent. When a tranquilizer is administered the barbiturate dosage can be reduced 10 to 25 percent.
- (3) Federal law restricts this drug to use by or on the order of a licensed veterinarian.

§ 522.2444b Sodium thiopental, sodium pentobarbital for injection.

(a) Specifications. Each gram of the drug contains 750 milligrams of sodium thiopental and 250 milligrams of sodium pentobarbital sterile powder for

dilution with sterile water for injection.

- (b) Sponsor. See No. 061133 in §510.600(c) of this chapter.
- (c) Conditions of use. (1) It is used as an anesthetic for intravenous administration to dogs and cats during short to moderately long surgical procedures.
 - (2) It is administered as follows:
- (i) For total anesthesia, it is given at approximately 10 to 12 milligrams per pound of body weight over a period of 3.5 to 5 minutes.
- (ii) When preanesthetic medication is used, it is important to wait at least an hour before administering thiopental and sodium pentobarbital for injection, and the dosage necessary for anesthesia is reduced. Usually ½ to ¾ the normal amount is adequate.
- (3) Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[40 FR 13858, Mar. 27, 1975, as amended at 47 FR 14149, Apr. 2, 1982; 66 FR 14073, Mar. 9, 2001]

§ 522.2470 Tiletamine hydrochloride and zolazepam hydrochloride for injection.

- (a) Specifications. Tiletamine hydrochloride and zolazepam hydrochloride for injection when reconstituted with sterile distilled water provides tiletamine hydrochloride and zolazepam hydrochloride equivalent to 50 milligrams of tiletamine base and 50 milligrams of zolazepam base per milliliter of solution.
- (b) Sponsor. See No. 000031 in \$510.600(c) of this chapter.
- (c) Conditions of use—(1) Indications for use. It is used for restraint or for anesthesia combined with muscle relaxation in cats and in dogs for restraint and minor procedures of short duration (30 minutes) requiring mild to moderate analgesia.
- (2) *Amount*. Expressed as milligrams of the drug combination:
- (i) In healthy dogs: An initial intramuscular dosage of 3 to 4.5 milligrams per pound of body weight for diagnostic purposes; 4.5 to 6 milligrams per pound of body weight for minor procedures of short duration such as